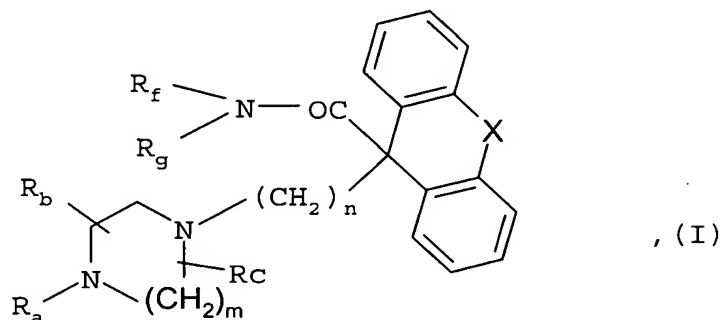


# LISTING OF CLAIMS

Claims 1-10 (Canceled)

Claim 11 (Currently Amended): A compound of the formula (I)



wherein

n denotes the number 1, 2, 3, 4 or 5,

m denotes the number 2 or 3,

X denotes a carbon-carbon bond, ~~an oxygen atom, a methylene, ethylene, imino or N-(C<sub>1-3</sub>-alkyl)-imino group,~~

R<sub>a</sub> denotes a phenyl group or a monocyclic heteroaryl group substituted by the groups R<sub>1</sub> and R<sub>2</sub>, wherein

R<sub>1</sub> denotes a hydrogen, fluorine, chlorine ~~or bromine atom~~, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms of the alkyl are optionally wholly or partly replaced by fluorine atoms, ~~a hydroxy group,~~ a C<sub>1-4</sub>-alkoxy group ~~wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms,~~ a phenoxy,

heteroaryloxy, phenyl-C<sub>1-3</sub>-alkoxy, carboxy, C<sub>1-3</sub>-alkoxycarbonyl, aminocarbonyl, C<sub>1-3</sub>-alkylaminocarbonyl, N,N-di-(C<sub>1-3</sub>-alkyl)-aminocarbonyl, nitro, or amino, C<sub>1-3</sub>-alkylamino, di-(C<sub>1-3</sub>-alkyl)-amino, phenyl-C<sub>1-3</sub>-alkyl-amino, N-(C<sub>1-3</sub>-alkyl)-phenyl-C<sub>1-3</sub>-alkylamino, C<sub>1-3</sub>-alkylcarbonylamino, N-(C<sub>1-3</sub>-alkyl)-C<sub>1-3</sub>-alkylcarbonylamino, C<sub>1-3</sub>-alkylsulphonylamino or N-(C<sub>1-3</sub>-alkyl)-C<sub>1-3</sub>-alkylsulphonylamino group, wherein the abovementioned phenyl of the phenoxy or heteroaryl moieties of the group R<sub>1</sub> are is optionally substituted by one to five fluorine, chlorine or methoxy bromine atoms, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a hydroxy group, or a C<sub>1-4</sub>-alkoxy group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, and

R<sub>2</sub> denotes a hydrogen, ~~fluorine, chlorine or bromine atom,~~ a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, ~~or a C<sub>1-4</sub>-alkoxy group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, or~~

R<sub>1</sub> and R<sub>2</sub> together represent a methylenedioxy group,

or R<sub>a</sub> denotes a monocyclic heteroaryl or phenyl group which is substituted in each case by a phenyl ~~or monocyclic heteroaryl~~ group, while the abovementioned phenyl groups and heteroaryl groups are optionally in each case substituted by a fluorine, chlorine or bromine atom, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, by a hydroxy, C<sub>1-3</sub>-alkoxy, carboxy, C<sub>1-3</sub>-alkoxycarbonyl, aminocarbonyl, C<sub>1-3</sub>-alkylaminocarbonyl or N,N-di-(C<sub>1-3</sub>-alkyl)-aminocarbonyl group,

R<sub>b</sub> and R<sub>c</sub> independently of one another denote a hydrogen atom or a C<sub>1-3</sub>-alkyl group and

R<sub>f</sub> and R<sub>g</sub>, which are identical or different, ~~denote~~ denotes hydrogen atom atoms, C<sub>1-6</sub>-alkyl groups wherein the hydrogen atoms of the alkyl are optionally wholly or partly

replaced by fluorine atoms, ~~C<sub>3-7</sub>-cycloalkyl groups, phenyl, heteroaryl, phenyl-C<sub>1-3</sub>-alkyl or heteroaryl-C<sub>1-3</sub>-alkyl groups, while wherein the abovementioned phenyl groups and heteroaryl groups are~~ optionally in each case be substituted by one to three fluorine ; chlorine or bromine atoms, by one to three C<sub>1-3</sub>-alkyl groups wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, by one to three hydroxy groups, one to three or C<sub>1-3</sub>-alkoxy groups wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, or by a carboxy, C<sub>1-3</sub>-alkoxycarbonyl, aminocarbonyl, C<sub>1-3</sub>-alkylaminocarbonyl, N,N-di-(C<sub>1-3</sub>-alkyl)-aminocarbonyl, N,N-di-(C<sub>1-3</sub>-alkyl)-amino, nitro or amino group, or

~~R<sub>f</sub> and R<sub>g</sub> together with the nitrogen atom between them denote a 3- to 7-membered cycloalkyleneimino group, while the methylene group in the 4 position of a 6- or 7-membered cycloalkyleneimino group is optionally replaced by an oxygen or sulphur atom, by a sulphinyl, sulphonyl, imino or N-(C<sub>1-3</sub>-alkyl)-imino group,~~

~~wherein the tricyclic group in the abovementioned formula I are mono- or disubstituted by fluorine or chlorine atoms, by methyl or methoxy groups and the substituents are identical or different,~~

~~and wherein the abovementioned heteroaryl groups in this claim are 6-membered heteroaryl groups containing one, two or three nitrogen atoms, or 5-membered heteroaryl groups containing one to four heteroatoms selected from nitrogen, oxygen and sulphur, while hydrogen atoms bound to nitrogen is optionally replaced by C<sub>1-3</sub>-alkyl groups,~~

R<sub>g</sub> is hydrogen;

or

the enantiomers, diastereomers isomers or the salts thereof.

Claim 12 (Currently amended): The compound according to claim 11, wherein

n denotes the number 3, 4 or 5 ;

m denotes the number 2 or 3,

X denotes a carbon-carbon bond, an oxygen atom, a methylene, ethylene, imino or N-(C<sub>1-3</sub>-alkyl)-imino group;

R<sub>a</sub> denotes a phenyl group or heteroaryl group substituted by the groups R<sub>1</sub> and R<sub>2</sub>, wherein

R<sub>1</sub> denotes a hydrogen, fluorine, chlorine or bromine atom, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a hydroxy group, a C<sub>1-4</sub>-alkoxy group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a phenoxy, heteroaryloxy, phenyl C<sub>1-3</sub>-alkoxy, carboxy, C<sub>1-3</sub>-alkoxycarbonyl, aminocarbonyl, C<sub>1-3</sub>-alkylaminocarbonyl, N,N-di-(C<sub>1-3</sub>-alkyl)-aminocarbonyl, nitro, amino, C<sub>1-3</sub>-alkylamino, di-(C<sub>1-3</sub>-alkyl)-amino, phenyl C<sub>1-3</sub>-alkyl-amino, N-(C<sub>1-3</sub>-alkyl)-phenyl C<sub>1-3</sub>-alkylamino, C<sub>1-3</sub>-alkylcarbonylamino, N-(C<sub>1-3</sub>-alkyl)-C<sub>1-3</sub>-alkylcarbonylamino, C<sub>1-3</sub>-alkylsulphonylamino or N-(C<sub>1-3</sub>-alkyl)-C<sub>1-3</sub>-alkylsulphonylamino group, wherein the abovementioned phenyl or heteroaryl moieties of the group R<sub>1</sub> are optionally substituted by one to five fluorine, chlorine or bromine atoms, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a hydroxy group, or a C<sub>1-4</sub>-alkoxy group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, and

R<sub>2</sub> denotes a hydrogen, fluorine, chlorine or bromine atom, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, or a C<sub>1-4</sub>-alkoxy group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, or

~~R<sub>1</sub> and R<sub>2</sub> together represent a methylenedioxy group,~~

~~or R<sub>a</sub> denotes a monocyclic heteroaryl or phenyl group which is substituted in each case by a phenyl or monocyclic heteroaryl group, wherein the abovementioned phenyl groups and heteroaryl groups are optionally in each case be substituted by a fluorine, chlorine or bromine atom, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, by a hydroxy or C<sub>1-3</sub>-alkoxy group,~~

~~R<sub>b</sub> and R<sub>c</sub> independently of one another denote a hydrogen atom or a C<sub>1-3</sub>-alkyl group and~~

~~R<sub>f</sub> and R<sub>g</sub>, which are identical or different, denote hydrogen atoms, C<sub>1-6</sub>-alkyl groups wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, C<sub>3-7</sub>-cycloalkyl groups, phenyl, heteroaryl, phenyl-C<sub>1-3</sub>-alkyl or heteroaryl-C<sub>1-3</sub>-alkyl groups, wherein the abovementioned phenyl groups and heteroaryl groups are optionally in each case be substituted by one to three fluorine, chlorine or bromine atoms, by one to three C<sub>1-3</sub>-alkyl groups wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, by one to three hydroxy groups, one to three C<sub>1-3</sub>-alkoxy groups wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, or by a carboxy, C<sub>1-3</sub>-alkoxycarbonyl, aminocarbonyl, C<sub>1-3</sub>-alkylaminocarbonyl, N,N-di-(C<sub>1-3</sub>-alkyl)-aminocarbonyl, N,N-di-(C<sub>1-3</sub>-alkyl)-amino, nitro or amino group, or and~~

~~R<sub>f</sub> and R<sub>g</sub> together with the nitrogen atom between them denote a 3- to 7-membered cycloalkyleneimino group, wherein the methylene group in the 4 position of a 6- or 7-membered cycloalkyleneimino group is optionally replaced by an oxygen or sulphur atom, by a sulphinyl, sulphonyl, imino or N-(C<sub>1-3</sub>-alkyl)-imino group .~~

Claim 13 (Currently amended): The compound according to claim 11, wherein

~~n denotes the number 3, 4 or 5,~~

m denotes the number 2 or 3,

X denotes a carbon-carbon bond or an oxygen atom,

R<sub>a</sub> denotes a phenyl group or heteroaryl group substituted by the groups R<sub>1</sub> and R<sub>2</sub>,  
wherein

R<sub>1</sub> denotes a hydrogen, fluorine, chlorine or bromine atom, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a hydroxy group, a C<sub>1-4</sub>-alkoxy group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a phenoxy, heteroaryloxy, phenyl-C<sub>1-3</sub>-alkoxy, carboxy, C<sub>1-3</sub>-alkoxycarbonyl, aminocarbonyl, C<sub>1-3</sub>-alkylaminocarbonyl, N,N-di-(C<sub>1-3</sub>-alkyl)-aminocarbonyl, nitro, amino, C<sub>1-3</sub>-alkylamino, di-(C<sub>1-3</sub>-alkyl)-amino, phenyl-C<sub>1-3</sub>-alkyl-amino, N-(C<sub>1-3</sub>-alkyl)-phenyl-C<sub>1-3</sub>-alkylamino, C<sub>1-3</sub>-alkylcarbonylamino, N-(C<sub>1-3</sub>-alkyl)-C<sub>1-3</sub>-alkylcarbonylamino, C<sub>1-3</sub>-alkylsulphonylamino or N-(C<sub>1-3</sub>-alkyl)-C<sub>1-3</sub>-alkylsulphonylamino group, wherein the abovementioned phenyl or heteroaryl moieties of the group R<sub>1</sub> are optionally substituted by one to five fluorine, chlorine or bromine atoms, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a hydroxy group, or a C<sub>1-4</sub>-alkoxy group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, and

R<sub>2</sub> denotes a hydrogen, fluorine, chlorine or bromine atom, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, or a C<sub>1-4</sub>-alkoxy group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, or

R<sub>1</sub> and R<sub>2</sub> together represent a methylenedioxy group,

~~or R<sub>a</sub> denotes a monocyclic heteroaryl or phenyl group which is substituted in each case by a phenyl or monocyclic heteroaryl group, wherein the abovementioned phenyl groups and heteroaryl groups are optionally in each case be substituted by a fluorine, chlorine or bromine atom, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, by a hydroxy or C<sub>1-3</sub>-alkoxy group,~~

R<sub>b</sub> and R<sub>c</sub> independently of one another denote a hydrogen atom or a methyl group and

~~R<sub>d</sub> denotes a hydrogen atom, a C<sub>1-6</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a C<sub>3-7</sub>-cycloalkyl group, phenyl, heteroaryl, phenyl-C<sub>1-3</sub>-alkyl or heteroaryl-C<sub>1-3</sub>-alkyl group, while the abovementioned phenyl groups and heteroaryl groups are optionally in each case be substituted by one to three fluorine, chlorine or bromine atoms, by one to three C<sub>1-3</sub>-alkyl groups wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, by one to three hydroxy groups, one to three C<sub>1-3</sub>-alkoxy groups wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, or by a nitro or amino group, and~~

~~R<sub>e</sub> denotes a hydrogen atom .~~

14( Currently amended): The compound according to claim 11, wherein

n denotes the number 4,

m denotes the number 2 ;

~~X denotes a carbon-carbon bond or an oxygen atom,~~

~~R<sub>a</sub> denotes a phenyl group or heteroaryl group substituted by the groups R<sub>1</sub> and R<sub>2</sub>, wherein~~

~~R<sub>1</sub> denotes a hydrogen, fluorine or chlorine atom, a C<sub>1-3</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a~~

~~C<sub>1-4</sub>-alkoxy group, a phenoxy group, a phenyl C<sub>1-3</sub>-alkoxy or a nitro or amino group;~~

~~wherein the abovementioned phenyl moiety of the phenoxy group is optionally substituted by a chlorine atom or by a methoxy group;~~

~~— R<sub>2</sub> denotes a hydrogen atom, a chlorine atom or a C<sub>1</sub>-C<sub>4</sub>-alkoxy group;~~

~~or R<sub>a</sub> denotes a monocyclic heteroaryl or phenyl group which is substituted in each case by a phenyl group;~~

~~R<sub>b</sub> and R<sub>c</sub> independently of one another denote a hydrogen atom or a C<sub>1-3</sub>-alkyl group and~~

~~R<sub>f</sub> denotes a C<sub>1</sub>-C<sub>6</sub>-alkyl group wherein the hydrogen atoms are optionally wholly or partly replaced by fluorine atoms, a phenyl C<sub>1-3</sub>-alkyl group, while the abovementioned phenyl group is optionally substituted in each case by a fluorine atom or by a C<sub>1</sub>-C<sub>3</sub>-alkoxy group, and~~

~~R<sub>g</sub> denotes a hydrogen atom .~~

Claim 15 (Currently amended): A compound chosen from

9-[4-(4-biphenyl-3-yl-piperazin-1-yl)-butyl]-9H-fluorene-9-carboxylic acid-(2,2,2-trifluoroethyl)-amide and

9-[4-(4-biphenyl-4-yl-piperazin-1-yl)-butyl]-9H-fluorene-9-carboxylic acid-(2,2,2-trifluoroethyl)-amide

or the enantiomers, diastereomers isomers and or the salts thereof.



Claim 16(Previously added): A physiologically acceptable salt of the compound according to claim 11.

Claim 17( Previously added): A pharmaceutical composition comprising a pharmaceutically effective amount of a compound according to claim 11 with one or more pharmaceutically acceptable inert carriers and/or diluents.

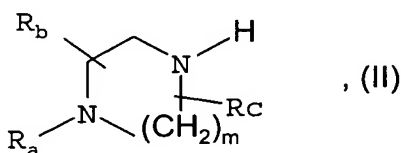
Claim 18 (Canceled).

Claim 19 (Previously added): A method of treating a disease selected from hyperlipidaemias, atherosclerosis and the clinical sequela thereof, diabetes mellitus, adiposity and pancreatitis, said method comprising administering to a patient in need thereof a pharmaceutically effective amount of a compound according to claim 11.

Claim 20 (Currently amended): The method according to ~~either of claims 18 or~~ claim 19 wherein the compound according to claim 11 is combined with another lipid-lowering agent.

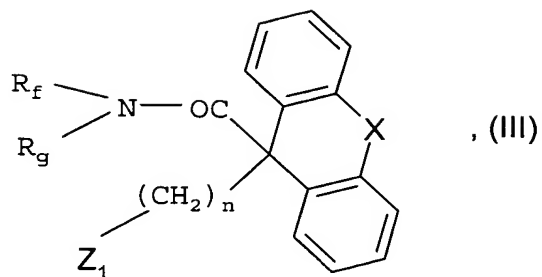
Claim 21( Currently amended): Process for preparing a compound of the formula (I) according to claim 11, comprising

a) reacting under suitable conditions a compound of formula



wherein

~~R<sub>a</sub>, R<sub>b</sub> and R<sub>c</sub> are defined as in claims 1,~~ with a compound of formula

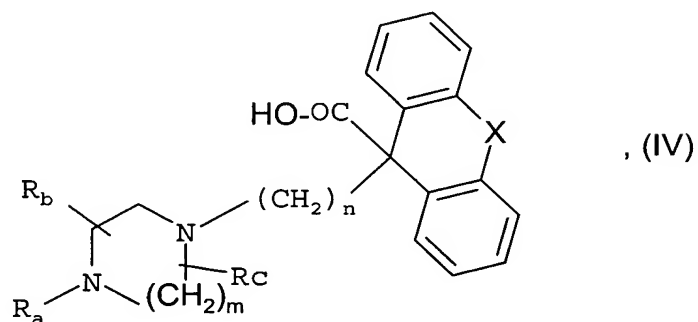


wherein

~~n, R<sub>f</sub>, R<sub>g</sub> and the tricyclic system are defined as in claims 1 and~~

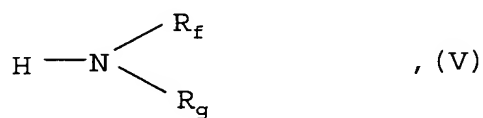
Z<sub>1</sub> denotes a nucleofugic leaving group, or

b) reacting under suitable conditions a compound of formula



wherein

~~the tricyclic system is defined as in claims 1, with an amine of formula~~



wherein

~~R<sub>f</sub> and R<sub>g</sub> are defined as in claims 1,~~ or with the reactive derivatives thereof and

c) optionally reducing under suitable conditions the product of a) or b) which contains a nitro group if desired into a corresponding amino compound and/or

d) if R<sub>f</sub> denotes a hydrogen atom alkylating under suitable conditions the product into a corresponding compound wherein R<sub>f</sub> denotes a ~~C<sub>1-3</sub>-alkyl~~ or phenyl-C<sub>1-3</sub>-alkyl group, and/or

e) cleaving under suitable conditions any protecting group using to protect reactive groups during the reactions and/or

resolving the product any of the product above into its stereoisomers and/or

converting any of the products above into the physiologically acceptable salts thereof.